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                 alerts (SDIs) affected
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                 alerts (SDIs) affected
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     12 DEC 17
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     13 DEC 17
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NEWS 14 DEC 30 EPFULL: New patent full text database to be available on STN
NEWS 15 DEC 30 CAPLUS - PATENT COVERAGE EXPANDED
NEWS 16 JAN 03 No connect-hour charges in EPFULL during January and
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L3 2 WEVLCWTWETCER/SQEP

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L11 ANSWER 1 OF 4 CAPLUS COPYRIGHT 2005 ACS on STN DUPLICATE 1

AN 2004:372929 CAPLUS

DN 140:395489

TI Sequences of blood-coagulation factor VIIa-binding peptides

IN Lazarus, Robert A.; Maun, Henry R.

PA Genentech, Inc., USA

SO U.S. Pat. Appl. Publ., 102 pp.

CODEN: USXXCO

DT Patent

LA English

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI US 2004087767	A1	20040506	US 2003-356257	20030130
PRAI US 2002-355420P	P	20020206		

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use); BIOL (Biological study); PREP (Preparation); USES (Uses)
        (factor VIIa-binding anticoagulant peptide sequence; sequences of
        blood-coagulation factor VIIa-binding peptides)
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     RL: PRP (Properties)
        (unclaimed sequence; sequences of blood-coagulation factor VIIa-binding
        peptides)
     This invention provides sequences of 6 blood-coagulation factor
AB
     VIIa-binding peptides. This invention provides novel compds. which
     prevent or block a FVIIa mediated or associated process or event such as the
     catalytic conversion of FX to FXa , FVII to FVIIa or FIX to FIXa.
     particular aspects, the compds. of the invention bind Factor VIIa (FVIIa
     ), its zymogen Factor VII (FVII). The invention also provides
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685513-39-7P

RL: BPN (Biosynthetic preparation); PRP (Properties); THU (Therapeutic

IT

358740-54-2P

685513-41-1P

685512-19-0P

685513-42-2P

685513-40-0P

pharmaceutical compns. comprising the novel compds. as well as their use in diagnostic, therapeutic, and prophylactic methods.

- L11 ANSWER 2 OF 4 CAPLUS COPYRIGHT 2005 ACS on STN
- AN 2003:443545 CAPLUS
- DN 139:159700
- TI Engineering Exosite Peptides for Complete Inhibition of Factor VIIa Using a Protease Switch with Substrate Phage
- AU Maun, Henry R.; Eigenbrot, Charles; Lazarus, Robert A.
- CS Department of Protein Engineering, Genentech, Inc., South San Francisco, CA, 94080, USA
- SO Journal of Biological Chemistry (2003), 278(24), 21823-21830 CODEN: JBCHA3; ISSN: 0021-9258
- PB American Society for Biochemistry and Molecular Biology
- DT Journal
- LA English
- IT 575431-91-3P, A 183X

RL: PAC (Pharmacological activity); PNU (Preparation, unclassified); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(engineering exosite peptides for complete inhibition of factor VIIa using a protease switch with substrate phage)

AB Limitations of current anticoagulant therapies have led us to develop two distinct classes of exosite peptide inhibitors for the initiator of the clotting process, the tissue factor-factor VIIa (TF·FVIIa) complex (Roberge, M., Santell, L., Dennis, M. S., Eigenbrot, C., Dwyer, M. A., and Lazarus, R. A. (2001) Biochem. 40, 9522-9531). Although both peptide classes are potent and selective inhibitors of TF·FVIIa, neither showed 100% inhibition at saturating concns. Crystal structures of these peptides in complex with the FVII/FVIIa protease domain revealed their distinct binding sites and close proximity to the active site. The favorable orientation of the 15-mer A-site peptide A-183 (EEWEVLCWTWETCER) suggested that a C-terminal extension into the FVIIa active site could yield a chimeric inhibitor that was not only potent and selective but complete as well. A novel two-step "protease switch" approach using substrate phage display was developed by first binding all phage containing A-183 and C-terminal extension libraries to immobilized and inactive FVIIa. Upon altering pH and adding TF to switch on FVIIa enzymic activity, only those phage released by proteolytic cleavage within the extension were propagated. This process selected for both preferred sequence and length in the extension, leading to a 27-mer peptide A-183X (EEWEVLCWTWETCERGEGVEEELWEWR) with a C-terminal 12-mer extension containing an Arg in the P1 position. A-183X was a more potent and complete inhibitor of FX activation, having a maximal extent of inhibition of .apprx.99% with an IC50 of 230 pM vs. A-183 which maximally inhibited to 74% with an IC50 of 1.5 nM. A-183X also had a maximal prolongation of the prothrombin time of 7.6- vs. 1.9-fold for A-183, making it a more effective anticoagulant.

RE.CNT 44 THERE ARE 44 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

- L11 ANSWER 3 OF 4 CAPLUS COPYRIGHT 2005 ACS on STN
- AN 2001:514493 CAPLUS
- DN 135:223287
- TI A novel exosite on coagulation factor VIIa and its molecular interactions with a new class of peptide inhibitors
- AU Roberge, Martin; Santell, Lydia; Dennis, Mark S.; Eigenbrot, Charles; Dwyer, Mary A.; Lazarus, Robert A.
- CS Department of Protein Engineering, Genentech Inc., South San Francisco, CA, 94080, USA
- SO Biochemistry (2001), 40(32), 9522-9531 CODEN: BICHAW; ISSN: 0006-2960
- PB American Chemical Society
- DT Journal
- LA English

ΙT 319927-97-4 325722-51-8 325722-64-3 358740-54-2 358740-54-2D, biotinylated derivs. RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); BIOL (Biological study) (protease domain exosite on coagulation factor VIIa and mol. interactions with A-series peptide inhibitors) A new inhibitory peptide binding exosite on the protease domain of AB coagulation Factor VIIa (FVIIa) has been identified. A novel series of peptide inhibitors of FVIIa, termed the "A-series" peptides, identified from peptide phage libraries and exemplified by peptide A-183, specifically bind at a site that is distinct from both the active site and the exosite of another recently described peptide inhibitor of FVIIa, E-76. Peptide A-183 prolonged TF-dependent clotting in human, but not rabbit plasma. Thus, a panel of human FVIIa mutants, containing 70 of the 76 rabbit sequence differences in the protease domain, localized the binding site to residues in the 60s loop and the C-terminus. The location of the exosite was refined by a series of FVIIa alanine mutants, which showed that proximal residues Trp 61 and Leu 251 were critical for binding. Kinetic and equilibrium binding consts. for zymogen FVII, FVIIa and TF FVIIa were determined using immobilized N-terminal biotinylated A-183 by surface plasmon resonance. No peptide binding to nine other human serine proteases was observed Key residues on the peptide were determined from binding to FVIIa and inhibition of FX activation using a series of alanine mutants of A-183 fused to the Z domain of protein A. Anal. of the mutagenesis data is presented in the context of a crystal structure of A-183 in complex with a version of zymogen FVII. The shape and proximity of this exosite to the active site may lend itself towards the design of new anticoagulants that inhibit FVIIa. RE.CNT 47 THERE ARE 47 CITED REFERENCES AVAILABLE FOR THIS RECORD . ALL CITATIONS AVAILABLE IN THE RE FORMAT L11 ANSWER 4 OF 4 CAPLUS COPYRIGHT 2005 ACS on STN AN 2001:496925 CAPLUS DN 135:221051 ΤI Selection and characterization of a new class of peptide exosite inhibitors of coaqulation factor VIIa ΑU Dennis, Mark S.; Roberge, Martin; Quan, Cliff; Lazarus, Robert A. CS Departments of Protein Engineering and Bioorganic Chemistry, Genentech Inc., South San Francisco, CA, 94080, USA SO Biochemistry (2001), 40(32), 9513-9521 CODEN: BICHAW; ISSN: 0006-2960 PB American Chemical Society DT Journal LA English IT 325722-64-3 358740-54-2 359635-57-7 325722-51-8 359635-58-8 359635-59-9 359635-60-2 359635-61-3 359635-62-4 359635-63-5 359635-64-6 359635-65-7 359635-67-9 359635-66-8 RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); PRP (Properties); BIOL (Biological study) (selection and characterization of peptide exosite inhibitors of coagulation factor VIIa) A new series of peptide inhibitors of human Factor VIIa (FVIIa) has been AB identified and affinity matured from naive and partially randomized peptide phage libraries selected against the immobilized tissue

AB A new series of peptide inhibitors of human Factor VIIa (FVIIa) has been identified and affinity matured from naive and partially randomized peptide phage libraries selected against the immobilized tissue factor Factor VIIa (TF·FVIIa) complex. These "A-series" peptides contain a single disulfide bond and a 13-residue minimal core required for maximal affinity. They are exemplified by peptide A-183 (EEWEVLCWTWETCER), which binds at a newly identified exosite on the FVIIa protease domain, described in the accompanying report [Roberge, M., Santell, L., Dennis, M. S., Eigenbrot, C., Dwyer, M. A., and Lazarus, R. A. (2001) Biochem. 40, XXXXXX-XXXXX]. A-183 was obtained from a trypsin digest of A-100-Z, a recombinant protein comprising A-183 and the Z domain

of protein A. Surprisingly, A-183 was a very potent inhibitor of TF·FVIIa, inhibiting activation of Factor X (FX) and Factor IX and amidolytic activity of Chromozym t-PA with IC50 values of 1.6 \pm 1.2, 3.5 ± 0.3 , and 8.5 ± 3.5 nM, resp. Kinetic anal. revealed that A-183 was a partial (hyperbolic) mixed-type inhibitor of FX activation having a Ki of 200 pM as well as a partial competitive inhibitor of amidolytic activity. The A-series peptides were also specific and potent inhibitors of TF-dependent clotting as measured in a prothrombin time (PT) clotting assay and had no effect on the TF-independent activated partial thromboplastin time. At saturating concns. of peptide, the maximal extent by which A-183 and A-100-Z inhibited the rate of FX activation was 78 \pm 3 and 89 \pm 6%, resp. The degree of inhibition of the rate of FX activation correlated with a maximum fold prolongation in the PT assay of 1.8-fold for A-183 and 3.3-fold for A-100-Z. The A-series peptides represent a new class of peptide exosite inhibitors that are capable of attenuating, rather than completely inhibiting, the activity of TF·FVIIa, potentially leading to anticoagulants with an increased therapeutic window.

RE.CNT 41 THERE ARE 41 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

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                alerts (SDIs) affected
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COST IN U.S. DOLLARS

ENTRY SESSION 211.39 211.60

FULL ESTIMATED COST

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L11 ANSWER 1 OF 4 CAPLUS COPYRIGHT 2005 ACS on STN DUPLICATE 1

AN 2004:372929 CAPLUS

DN 140:395489

TI Sequences of blood-coagulation factor VIIa-binding peptides

IN Lazarus, Robert A.; Maun, Henry R.

PA Genentech, Inc., USA

SO U.S. Pat. Appl. Publ., 102 pp.

CODEN: USXXCO

DT Patent

LA English

FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI US 2004087767	A1	20040506	US 2003-356257	20030130
PRAT US 2002-355420P	P	20020206		

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(factor VIIa-binding anticoagulant peptide sequence; sequences of
        blood-coagulation factor VIIa-binding peptides)
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     RL: PRP (Properties)
        (unclaimed sequence; sequences of blood-coagulation factor VIIa-binding
        peptides)
AB
     This invention provides sequences of 6 blood-coagulation factor
     VIIa-binding peptides. This invention provides novel compds. which
     prevent or block a FVIIa mediated or associated process or event such as the
     catalytic conversion of FX to FXa , FVII to FVIIa or FIX to FIXa.
     particular aspects, the compds. of the invention bind Factor VIIa (FVIIa
     ), its zymogen Factor VII (FVII). The invention also provides
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685513-39-7P

RL: BPN (Biosynthetic preparation); PRP (Properties); THU (Therapeutic

use); BIOL (Biological study); PREP (Preparation); USES (Uses)

685513-40-0P

IT

358740-54-2P

685513-41-1P

685512-19-0P

685513-42-2P

pharmaceutical compns. comprising the novel compds. as well as their use in diagnostic, therapeutic, and prophylactic methods.

- L11 ANSWER 2 OF 4 CAPLUS COPYRIGHT 2005 ACS on STN
- AN 2003:443545 CAPLUS
- DN 139:159700
- TI Engineering Exosite Peptides for Complete Inhibition of Factor VIIa Using a Protease Switch with Substrate Phage
- AU Maun, Henry R.; Eigenbrot, Charles; Lazarus, Robert A.
- CS Department of Protein Engineering, Genentech, Inc., South San Francisco, CA, 94080, USA
- SO Journal of Biological Chemistry (2003), 278(24), 21823-21830 CODEN: JBCHA3; ISSN: 0021-9258
- PB American Society for Biochemistry and Molecular Biology
- DT Journal
- LA English
- IT 575431-91-3P, A 183X

RL: PAC (Pharmacological activity); PNU (Preparation, unclassified); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(engineering exosite peptides for complete inhibition of factor VIIa using a protease switch with substrate phage)

AB Limitations of current anticoagulant therapies have led us to develop two distinct classes of exosite peptide inhibitors for the initiator of the clotting process, the tissue factor-factor VIIa (TF·FVIIa) complex (Roberge, M., Santell, L., Dennis, M. S., Eigenbrot, C., Dwyer, M. A., and Lazarus, R. A. (2001) Biochem. 40, 9522-9531). Although both peptide classes are potent and selective inhibitors of TF·FVIIa, neither showed 100% inhibition at saturating concns. Crystal structures of these peptides in complex with the FVII/FVIIa protease domain revealed their distinct binding sites and close proximity to the active site. favorable orientation of the 15-mer A-site peptide A-183 (EEWEVLCWTWETCER) suggested that a C-terminal extension into the FVIIa active site could yield a chimeric inhibitor that was not only potent and selective but complete as well. A novel two-step "protease switch" approach using substrate phage display was developed by first binding all phage containing A-183 and C-terminal extension libraries to immobilized and inactive FVIIa. Upon altering pH and adding TF to switch on FVIIa enzymic activity, only those phage released by proteolytic cleavage within the extension were propagated. This process selected for both preferred sequence and length in the extension, leading to a 27-mer peptide A-183X (EEWEVLCWTWETCERGEGVEEELWEWR) with a C-terminal 12-mer extension containing an Arg in the P1 position. A-183X was a more potent and complete inhibitor of FX activation, having a maximal extent of inhibition of .apprx.99% with an IC50 of 230 pM vs. A-183 which maximally inhibited to 74% with an IC50 of 1.5 nM. A-183X also had a maximal prolongation of the prothrombin time of 7.6- vs. 1.9-fold for A-183, making it a more effective anticoagulant.

RE.CNT 44 THERE ARE 44 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

- L11 ANSWER 3 OF 4 CAPLUS COPYRIGHT 2005 ACS on STN
- AN 2001:514493 CAPLUS
- DN 135:223287
- TI A novel exosite on coagulation factor VIIa and its molecular interactions with a new class of peptide inhibitors
- AU Roberge, Martin; Santell, Lydia; Dennis, Mark S.; Eigenbrot, Charles; Dwyer, Mary A.; Lazarus, Robert A.
- CS Department of Protein Engineering, Genentech Inc., South San Francisco, CA, 94080, USA
- SO Biochemistry (2001), 40(32), 9522-9531 CODEN: BICHAW; ISSN: 0006-2960
- PB American Chemical Society
- DT Journal
- LA English

319927-97-4 325722-51-8 325722-64-3 358740-54-2 IT 358740-54-2D, biotinylated derivs. RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); BIOL (Biological study) (protease domain exosite on coagulation factor VIIa and mol. interactions with A-series peptide inhibitors) A new inhibitory peptide binding exosite on the protease domain of AB coagulation Factor VIIa (FVIIa) has been identified. A novel series of peptide inhibitors of FVIIa, termed the "A-series" peptides, identified from peptide phage libraries and exemplified by peptide A-183, specifically bind at a site that is distinct from both the active site and the exosite of another recently described peptide inhibitor of FVIIa, E-76. Peptide A-183 prolonged TF-dependent clotting in human, but not rabbit plasma. Thus, a panel of human FVIIa mutants, containing 70 of the 76 rabbit sequence differences in the protease domain, localized the binding site to residues in the 60s loop and the C-terminus. The location of the exosite was refined by a series of FVIIa alanine mutants, which showed that proximal residues Trp 61 and Leu 251 were critical for binding. and equilibrium binding consts. for zymogen FVII, FVIIa and TF·FVIIa

plasmon resonance. No peptide binding to nine other human serine proteases was observed Key residues on the peptide were determined from binding

to FVIIa and inhibition of FX activation using a series of alanine mutants of A-183 fused to the Z domain of protein A. Anal. of the mutagenesis data is presented in the context of a crystal structure of A-183 in complex with a version of zymogen FVII. The shape and proximity of this exosite to the active site may lend itself towards the design of new anticoagulants that inhibit FVIIa.

were determined using immobilized N-terminal biotinylated A-183 by surface

RE.CNT 47 THERE ARE 47 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L11 ANSWER 4 OF 4 CAPLUS COPYRIGHT 2005 ACS on STN

AN 2001:496925 CAPLUS

DN 135:221051

TI Selection and characterization of a new class of peptide exosite inhibitors of coagulation factor VIIa

AU Dennis, Mark S.; Roberge, Martin; Quan, Cliff; Lazarus, Robert A.

CS Departments of Protein Engineering and Bioorganic Chemistry, Genentech Inc., South San Francisco, CA, 94080, USA

SO Biochemistry (2001), 40(32), 9513-9521 CODEN: BICHAW; ISSN: 0006-2960

PB American Chemical Society

DT Journal

LA English

IT 325722-51-8 325722-64-3 358740-54-2 359635-57-7

359635-58-8 359635-59-9 359635-60-2

359635-61-3 359635-62-4 359635-63-5 359635-64-6 359635-65-7

359635-66-8 359635-67-9

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); PRP (Properties); BIOL (Biological study)

(selection and characterization of peptide exosite inhibitors of coagulation factor VIIa)

AB A new series of peptide inhibitors of human Factor VIIa (FVIIa) has been identified and affinity matured from naive and partially randomized peptide phage libraries selected against the immobilized tissue factor Factor VIIa (TF·FVIIa) complex. These "A-series" peptides contain a single disulfide bond and a 13-residue minimal core required for maximal affinity. They are exemplified by peptide A-183 (EEWEVLCWTWETCER), which binds at a newly identified exosite on the FVIIa protease domain, described in the accompanying report [Roberge, M., Santell, L., Dennis, M. S., Eigenbrot, C., Dwyer, M. A., and Lazarus, R. A. (2001) Biochem. 40, XXXXXX-XXXXX]. A-183 was obtained from a trypsin digest of A-100-Z, a recombinant protein comprising A-183 and the Z domain

of protein A. Surprisingly, A-183 was a very potent inhibitor of TF·FVIIa, inhibiting activation of Factor X (FX) and Factor IX and amidolytic activity of Chromozym t-PA with IC50 values of 1.6 \pm 1.2, 3.5 \pm 0.3, and 8.5 \pm 3.5 nM, resp. Kinetic anal. revealed that A-183 was a partial (hyperbolic) mixed-type inhibitor of FX activation having a Ki of 200 pM as well as a partial competitive inhibitor of amidolytic activity. The A-series peptides were also specific and potent inhibitors of TF-dependent clotting as measured in a prothrombin time (PT) clotting assay and had no effect on the TF-independent activated partial thromboplastin time. At saturating concns. of peptide, the maximal extent by which A-183 and A-100-Z inhibited the rate of FX activation was 78 \pm 3 and 89 ± 6%, resp. The degree of inhibition of the rate of FX activation correlated with a maximum fold prolongation in the PT assay of 1.8-fold for A-183 and 3.3-fold for A-100-Z. The A-series peptides represent a new class of peptide exosite inhibitors that are capable of attenuating, rather than completely inhibiting, the activity of TF·FVIIa, potentially leading to anticoagulants with an increased therapeutic window.

RE.CNT 41 THERE ARE 41 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

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L13 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2005 ACS on STN

325722-42-7

RL: BAC (Biological activity or effector, except adverse); BPR (Biological process); BSU (Biological study, unclassified); PRP (Properties); BIOL (Biological study); PROC (Process)

(factor VIIa antagonists for diagnostic or therapeutic use)

=> d 113 hit bib 1-2

L13 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2005 ACS on STN

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RL: BAC (Biological activity or effector, except adverse); BPR (Biological
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        (factor VIIa antagonists for diagnostic or therapeutic use)
     2001:115174 CAPLUS
AN
DN
     134:168300
     Factor VIIa antagonists for diagnostic or therapeutic use
ΤI
IN
     Dennis, Mark S.
PΑ
     Genentech, Inc., USA
SO
     PCT Int. Appl., 80 pp.
     CODEN: PIXXD2
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     English
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       Genentech, Inc., South San Francisco, CA (U.S. corporation)
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       US 2003-639076 .
                          A1
                               20030811 (10)
       Continuation of Ser. No. US 2000-632429, filed on 4 Aug 2000, PENDING
RLI
PRAI
       US 1999-147627P
                           19990806 (60)
       US 1999-150315P
                           19990823 (60)
DT
       Utility
FS
       APPLICATION
       GENENTECH, INC., 1 DNA WAY, SOUTH SAN FRANCISCO, CA, 94080
LREP
CLMN
       Number of Claims: 31
ECL
       Exemplary Claim: 1
DRWN
       4 Drawing Page(s)
LN.CNT 2987
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
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ΙT

325722-42-7

COST IN U.S. DOLLARS SINCE FILE TOTAL

FULL ESTIMATED COST ENTRY SESSION 27.77 239.37

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS) SINCE FILE TOTAL

CA SUBSCRIBER PRICE ENTRY SESSION
-2.92 -2.92

FILE 'REGISTRY' ENTERED AT 15:12:05 ON 04 JAN 2005 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS. COPYRIGHT (C) 2005 American Chemical Society (ACS)

Property values tagged with IC are from the ZIC/VINITI data file provided by InfoChem.

STRUCTURE FILE UPDATES: 3 JAN 2005 HIGHEST RN 807382-78-1 DICTIONARY FILE UPDATES: 3 JAN 2005 HIGHEST RN 807382-78-1

TSCA INFORMATION NOW CURRENT THROUGH MAY 21, 2004

Please note that search-term pricing does apply when conducting SmartSELECT searches.

Crossover limits have been increased. See HELP CROSSOVER for details.

Experimental and calculated property data are now available. For more information enter HELP PROP at an arrow prompt in the file or refer to the file summary sheet on the web at: http://www.cas.org/ONLINE/DBSS/registryss.html

Claim

=> FIL CAPLUS BIOSIS MEDLINE PCTFULL USPATFULL EUROPATFULL JAPIO SCISEARCH EMBASE COST IN U.S. DOLLARS SINCE FILE TOTAL

FULL ESTIMATED COST ENTRY SESSION 58.87 298.24

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS) SINCE FILE TOTAL

CA SUBSCRIBER PRICE

ENTRY SESSION

0.00 -2.92

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FILE 'PCTFULL' ENTERED AT 15:17:14 ON 04 JAN 2005 COPYRIGHT (C) 2005 Univentio

FILE 'USPATFULL' ENTERED AT 15:17:14 ON 04 JAN 2005 CA INDEXING COPYRIGHT (C) 2005 AMERICAN CHEMICAL SOCIETY (ACS)

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=> FIL REGISTRY COST IN U.S. DOLLARS SINCE FILE TOTAL ENTRY SESSION FULL ESTIMATED COST 9.03 307.27 DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS) SINCE FILE TOTAL ENTRY SESSION CA SUBSCRIBER PRICE 0.00 -2.92

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Property values tagged with IC are from the ZIC/VINITI data file provided by InfoChem.

STRUCTURE FILE UPDATES: 3 JAN 2005 HIGHEST RN 807382-78-1 DICTIONARY FILE UPDATES: 3 JAN 2005 HIGHEST RN 807382-78-1

TSCA INFORMATION NOW CURRENT THROUGH MAY 21, 2004

Please note that search-term pricing does apply when conducting SmartSELECT searches.

Crossover limits have been increased. See HELP CROSSOVER for details.

Experimental and calculated property data are now available. For more information enter HELP PROP at an arrow prompt in the file or refer to the file summary sheet on the web at: http://www.cas.org/ONLINE/DBSS/registryss.html

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L11 4 DUP REM L10 (1 DUPLICATE REMOVED)

L12 7 S L2

L13 2 S L12 NOT L10

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24 S [WFL] E [LV] [LIMV] C [WFLM] TWETCE [RKLW] / SQSP

L15 22 S [WFLA] E [VIA] LC [WFLMA] TWETCER/SQSP

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=> s l14 or l15

L14

L16 28 L14 OR L15

=> FIL CAPLUS BIOSIS MEDLINE PCTFULL USPATFULL EUROPATFULL JAPIO SCISEARCH EMBASE USPAT2 EUROPATFULL

COST IN U.S. DOLLARS

SINCE FILE TOTAL ENTRY SESSION

FULL ESTIMATED COST

0.43 307.70

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)

SINCE FILE TOTAL ENTRY SESSION

CA SUBSCRIBER PRICE

0.00 -2.92

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FILE 'USPATFULL' ENTERED AT 15:18:15 ON 04 JAN 2005
CA INDEXING COPYRIGHT (C) 2005 AMERICAN CHEMICAL SOCIETY (ACS).

'EUROPATFULL' IS NOT A VALID FILE NAME

FILE 'JAPIO' ENTERED AT 15:18:15 ON 04 JAN 2005 COPYRIGHT (C) 2005 Japanese Patent Office (JPO) - JAPIO

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L17 7 L14

^{=&}gt; s 114

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L23
    2004 372929 CAPLUS
AN
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DN
     Sequences of blood-coagulation factor VIIa-binding peptides
ΤI
     Lazarus, Robert A.; Maun, Henry R.
IN
     Genentech, Inc., USA
PA
     U.S. Pat. Appl. Publ., 102 pp.
SO
     CODEN: USXXCO
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FAN.CNT 1
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                                DATE
                                            APPLICATION NO.
                                                                   DATE
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    ANSWER 2 OF 6 USPATFULL on STN
L23
       2004:101678 USPATFULL
       FVIIa antagonists
       Dennis, Mark S., San Carlos, CA/ UNITED STATES
       Genentech, Inc., South San Trancisco, CA (U.S. corporation)
                               200404/22
       US 2004077547
                          Αl
                               20030 (10)
       US 2003-639076
                          Α1
       Continuation of Ser. No. U$ 2000-632429, filed on 4 Aug 2000, PENDING
                           19990808 (60)
       US 1999-147627P
       US 1999-150315P
                           19990823 (60)
       Utility
       APPLICATION
       GENENTECH, INC., 1 DNA WAY, SOUTH SAN FRANCISCO, CA, 94080
      Number of Claims: 31
       Exemplary Claim: 1
       4 Drawing Page(s)
LN.CNT 2987
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
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ANΤI

IN

PA

PΙ

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RLI

DT

FS

LREP CLMN

ECL

DRWN

PRAI

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IT 325722-42-7
        (factor VIIa antagonists for diagnostic or therapeutic use)
L23
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AN
     2003:443545 CAPLUS
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     139:159700
ΤI
     Engineering Exosite Peptides for Complete Inhibition of Factor VIIa Using
     a Protease Switch with Substrate Phage
     Maun, Henry R.; Eigenbrot, Charles; Lazarus, Robert A.
AU
     Department of Protein Engineering, Genentech, Inc., South San Francisco,
CS
     CA, 94080, USA
                                     (2003), 278(24), 21823-21830
     Journal of Biological Chemistry
SO
     CODEN: JBCHA3; ISSN: 0021-9258
PB
     American Society for Biochemistry and Molecular Biology
DT
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LA
     English
RE.CNT 44
              THERE ARE 44 CITED REFERENCES AVAILABLE FOR THIS RECORD
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     134:168300
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     Dennis, Mark S.
PΑ
     Genentech, Inc., USA
SO
     PCT Int. Appl., 80 pp.
     CODEN: PIXXD2
DT
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LA
     English
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OS
IT
     325722-42-7
     RL: BAC (Biological activity or effector, except adverse); BPR (Biological
     process); BSU (Biological study, unclassified); PRP (Properties); BIOL
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(factor VIIa antagonists for diagnostic or therapeutic use) ANSWER 5 OF 6 CAPLUS COPYRIGHT 2005 ACS on STN 2001:514493 CAPLUS AN DN 135:223287 A novel exosite on coagulation factor VIIa and its molecular interactions TI with a new class of peptide inhibitors AU Roberge, Martin; Santell, Lydia; Dennis, Mark S.; Eigenbrot, Charles; Dwyer, Mary A.; Lafarus, Robert A. Department of Provein Engineering, Genentech Inc., South San Francisco, CS CA, 94080, USA | Biochemistry (2001), 40(32), 9522-9531 CODEN: BICHAW; USA: 0006-2960 SO PBAmerican Chemical Society DTJournal English LA RE.CNT 47 THERE ARE 47 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT TT 319927-97-4 325722-51-8 325722-64-3 **358740-54-2** 358740-54-2D, biotinylated derivs. RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); BIOL (Biological study) (protease domain exosite on coagulation factor VIIa and mol. interactions with A-series peptide inhibitors) ANSWER 6 OF 6 CAPLUS COPYRIGHT 2005 ACS on STN T₁2.3 AN 2001:496925 CAPLUS DN 135:221051 ΤI Selection and characterization of a new class of peptide exosite inhibitors of coagulation factor VIIa Dennis, Mark S.; Roberge, Martin; Quan, Cliff; Lazarus, Robert A. Departments of Protein Engineering and Bioorganic Chemistry, Genentech ΔΙΙ CS Inc., South San Francisco, CA, 94080, USA Biochemistry (2001), 40(32), 9513-9521 CODEN: BICHAW; USAN: 0006-2960 SO American Chemical Society PB DT Journal LA English THERE ARE 41 CITED REFERENCES AVAILABLE FOR THIS RECORD RE.CNT 41 ALL CITATIONS AVAILABLE IN THE RE FORMAT IT 325722-51-8 325722-64-3 **358740-54-2 359635-57-7** 359635-58-8 359635-59-9 359635-60-2 359635-61-3 359635-62-4 359635-63-5 359635-64-6 359635-67-9 359635-65-7 359635-66-8 RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); PRP (Properties); BIOL (Biological study) (selection and characterization of peptide exosite inhibitors of coagulation factor VIIa) => FIL REGISTRY COST IN U.S. DOLLARS SINCE FILE TOTAL ENTRY SESSION FULL ESTIMATED COST 36.80 344.50 DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS) SINCE FILE TOTAL ENTRY SESSION CA SUBSCRIBER PRICE 0.00 -2.92

(Biological study); PROC (Process)

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Experimental and calculated property data are now available. For more information enter HELP PROP at an arrow prompt in the file or refer to the file summary sheet on the web at: http://www.cas.org/ONLINE/DBSS/registryss.html

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PB Kazusa DNA Research Institute DT Journal LΑ English THERE ARE 30 CITED REFERENCES AVAILABLE FOR THIS RECORD RE.CNT 30 ALL CITATIONS AVAILABLE IN THE RE FORMAT => d 130 bib hit ANSWER 1 OF 224 CAPLUS COPYRIGHT 2005 ACS on STN L30 2004:1012134 CAPLUS AN DN 141:421056 Expressed sequence tags and encoded human proteins ΤI TN Edwards, Jean-Baptiste Dumas Milne; Duclert, Aymeric; Giordano, Jean-Yves PΑ Genset S.A., Fr. U.S., 72 pp., Cont.-in-part of Appl. No. PCT/IB99/00712. SO CODEN: USXXAM DT Patent LΑ English FAN.CNT 2 PATENT NO. KIND DATE APPLICATION NO. DATE PΙ US 6822072 B1 20041123 US 1999-471276 19991221 WO 9953051 19991021 WO 1999-IB712 19990409 <--A2 A3 WO 9953051 20000406 W: AU, CA, JP, US RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE PRAI US 1998-57719 B2 19980409 · B2 US 1998-69047 19980428 WO 1999-IB712 A2 19990409 THERE ARE 22 CITED REFERENCES AVAILABLE FOR THIS RECORD RE.CNT 22 ALL CITATIONS AVAILABLE IN THE RE FORMAT APPLICATION NO. PATENT NO. KIND DATE DATE _ _ _ _ _____ ----------20041123 US 1999-471276 PΤ US 6822072 В1 19991221 WO 9953051 WO 1999-IB712 19990409 <--A2 19991021 WO 9953051 Α3 20000406 W: AU, CA, JP, US RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE IT 246537-95-1 246537-99-5 220593-06-6 225514-15-8 246538-06-7 246538-07-8 246538-08-9 246538-09-0 246538-10-3 246538-24-9 246538-74-9 246538-77-2 246877-53-2 246877-54-3 246877-55-4 246877-58-7 **246877-59-8** 246877-56-5 246877-57-6 246877-64-5 246877-60-1 246877-61-2 246877-62-3 246877-63-4 246877-65-6 246877-66-7 246877-67-8 246877-68-9 246877-69-0 246877-70-3 246877-71-4 246877-72-5 246877-73-6 246877-74-7 246877-75-8 246877-76-9 246877-77-0 246877-78-1 246877-79-2 246877-80-5 246877-81-6 246877-82-7 246877-83-8 246877-84-9 246877-85-0 246877-86-1 246877-87-2 246877-88-3 246877-89-4 246877-90-7 246877-91-8 246877-92-9 246877-93-0 246877-94-1 246877-95-2 246877-96-3 246877-97-4 246877-98-5 246877-99-6 246878-00-2 246878-01-3 246878-02-4 246878-03-5 246878-04-6 246878-05-7 246878-06-8 246878-07-9 246878-08-0 246878-09-1 246878-10-4 246878-11-5 246878-12-6 246878-13-7 246878-14-8 246878-15-9 246878-16-0 246878-17-1 246878-18-2 246878-19-3 246878-20-6 246878-21-7 246878-22-8 246878-23-9 246878-24-0 246878-25-1 246878-26-2 246878-27-3 246878-28-4 246878-29-5 246878-30-8 246878-31-9 246878-32-0 246878-33-1 246878-34-2

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     Cloning and cDNA and deduced amino acid sequences of 97 human secreted
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    Ruben, Steven M.; Florence, Kimberly A.; Ni, Jian; Rosen, Craig A.;
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     Paul E.; Wei, Ying-fei; Brewer, Laurie A.; Soppet, Daniel R.; Lafleur,
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    U.S. Pat. Appl. Publ., 453 pp., Cont.-in-part of U.S. Ser. No. 892,877.
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    Mammalian cytokine AK155 polypeptides, polynucleotides and antibodies for
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    De Waal, Malefyt Rene; Flickensher, Helmut; Fleckenstein, Bernhard;
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       Morris, Jill A., Chalfont, PA, United States
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       Pavan, William J., Derwood, MD, United States
       Ashlock, Melissa A., Mont Vernon, NH, United States
       Loftus, Stacie K., Burtonsville, MD, United States
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TI
    Protein and cDNA sequences encoding a human IL-10 homolog, designated
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IN
    Knappe, Andrea; Fickenscher, Helmut; Fleckenstein, Bernhard
PΑ
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SO
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TT
    Cascieri, Margaret A.; MacNeil, Douglas J.; Shiao, Lin-lin; Weinberg,
IN
    David H.; Tan, Carina P.; Linemeyer, David L.; Strader, Catherine D.
PA
    Merck and Co., Inc., USA
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     Mammalian sperm protein pkdrej, its cDNA and methods of identifying
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IN
     Harris, Peter Charles; Hugues, James Raymond; Ward, Christopher James
PA
     Medical Research Council, UK
     PCT Int. Appl., 41 pp.
SO
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     Cloning and cDNA and deduced amino acid sequences of 97 human secreted
IN
     Ruben, Steven M.; Florence, Kimberly; Ni, Jian; Rosen, Craig A.; Carter,
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     E.; Wei, Fing-Fei; Brewer, Laurie A.; Soppet, Daniel R.; Lafleur, David
     W.; Endress, Gregory A.; Ebner, Reinhard
PΑ
     Human Genome Sciences, Inc., USA
     PCT Int. Appl., 475 pp.
SO
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AN
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ТT
     Human receptor (REC) polypeptides and polynucleotides, sequences, and
     biological and therapeutic uses thereof
TN
     Hillman, Jennifer L.; Bandman, Olga; Tang, Y. Tom; Yue, Henry; Lal,
     Preeti; Corley, Neil C.; Guegler, Karl J.; Patterson, Chandra
PA
     Incyte Pharmaceuticals, Inc., USA
SO
     PCT Int. Appl., 94 pp.
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DT
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IN
    Dumas Milne Edwards, Jean-Baptiste; Duclert, Aymeric; Giordano, Jean-Yves
PA
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    PCT Int. Appl., 837 pp.
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     Assays for screening compounds which interact with cation channel
ΤI
     proteins, mutant prokaryotic cation channel proteins, and uses thereof
TN
     MacKinnon, Roderick
PA
     The Rockefeller University, USA
so
     PCT Int. Appl., 165 pp.
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LA
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FAN.CNT 2
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     WO 9947923
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     WO 1999-US6307
                                19990322
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     ANSWER 16 OF 224 CAPLUS COPYRIGHT 2005 ACS on STN
L30
     1999:614169 CAPLUS
AN
DN
     131:238829
     Human and murine G protein-coupled heptahelical receptor D6 and its cDNA
TI
     sequences and therapeutic uses
     Graham, Gerard J.; Benjamin, Nibbs Robert J.; Gonzalo, Jose-Angel;
IN
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Gutierrez-Ramos, Jose-Carlos

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SO
     PCT Int. Appl., 152 pp.
     CODEN: PIXXD2
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RE.CNT 4
             THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD
             ALL CITATIONS AVAILABLE IN THE RE FORMAT
L30
    ANSWER 17 OF 224 CAPLUS COPYRIGHT 2005 ACS on STN
AN
    1999:576793 CAPLUS
DN
    131:195462
    Protein and cDNA sequences for human and mouse IL-9 Induced Calcium
TI
    Activated Chloride Channels (ICACC) and uses thereof in the treatment of
     atopic allergies, asthma, inflammatory bowel disease, and cystic fibrosis
IN
    Holroyd, Kenneth J.; Levitt, Roy C.; Maloy, W. Lee; Louahed, Jamila;
    McLane, Mike; Nicolaides, Nicholas C.; Zhou, Yuhong; Dong, Qu
PA
    Magainin Pharmaceuticals, Inc., USA
    PCT Int. Appl., 75 pp.
SO
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Millennium Pharmaceuticals, Inc., USA; CRC Technology Limited

PA

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               THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD
               ALL CITATIONS AVAILABLE IN THE RE FORMAT
L30
     ANSWER 18 OF 224 CAPLUS COPYRIGHT 2005 ACS on STN
     1999:388051 CAPLUS
AN
DN
     131:56596
ΤI
     Phaseolus genes expressed during senescence and their promoters and the
     stage-specific expression of foreign genes
     Gepstein, Shimon; Hajuoje, Taleb; Rosner, Amalia
IN
     Vitality Biotechnologies, Inc., USA
PA
so
     PCT Int. Appl., 69 pp.
     CODEN: PIXXD2
DT
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PRAI US 1997-67898P
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              THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD
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               ALL CITATIONS AVAILABLE IN THE RE FORMAT
     ANSWER 19 OF 224 CAPLUS COPYRIGHT 2005 ACS on STN
L30
     1999:48801 CAPLUS
AN
DN
     130:120476
ΤI
     Genes for Niemann-Pick type C disease
     Carstea, Eugene D.; Tagle, Danilo A.; Morris, Jill A.; Pentchev, Peter G.;
IN
     Pavan, William J.; Rosenfeld, Melissa A.; Loftus, Stacie K.; Gu, Jessie
     United States Dept. of Health and Human Services, USA
PΑ
SO
     PCT Int. Appl., 100 pp.
     CODEN: PIXXD2
DT
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LΑ
     English
FAN.CNT 1
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     AU 9882869
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                                              US 2002-208731
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PRAI US 1997-51682P
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     WU 1998-US13862 W
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RE.CNT 9
             THERE ARE 9 CITED REFERENCES AVAILABLE FOR THIS RECORD
             ALL CITATIONS AVAILABLE IN THE RE FORMAT
L30 ANSWER 20 OF 224 CAPLUS COPYRIGHT 2005 ACS on STN
AN
     1999:802470 CAPLUS
DN
     132:45817
ΤI
     Cloning of gene for cytochrome bd type quinol oxidase from Brevibacterium
     lactofermentum
IN
     Sone, Nobufumi
PΑ
     Ajinomoto Co., Inc., Japan
SO
     Jpn. Kokai Tokkyo Koho, 19 pp.
     CODEN: JKXXAF
DT
     Patent
LA
     Japanese
FAN.CNT 1
     PATENT NO.
                      KIND DATE
                                        APPLICATION NO.
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A3
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                        Α
=> d 130 21-30 Ti in so pn
L30
    ANSWER 21 OF 224 USPATFULL on STN
ΤI
       Proteins involved in the synthesis and assembly of O-antigen in
       Pseudomonas aeruginosa
IN
      Lam, Joseph S., Guelph, Canada
      Burrows, Lori, Guelph, Canada
       Charter, Deborah, Guelph, Canada
       de Kievit, Teresa, Guelph, Canada
PΙ
      US 5994072
                             19991130
                                                                 <--
    ANSWER 22 OF 224 USPATFULL on STN
L30
      Rolling mill roll stand
ΤI
IN
       Woodrow, Harold E., Northboro, MA, United States
       Shore, T. Michael, Princeton, MA, United States
PΙ
      US 5983694
                             19991116
    ANSWER 23 OF 224 USPATFULL on STN
L30
      Fine magnetic particles containing useful proteins bound thereto,
TI
      process for producing the same, and use thereof
      Matsunaga, Tadashi, B-506, 2-40, Saiwai-cho, Funchi-shi, Tokyo, 183,
IN
       Japan
      Kamiya, Shinji, Tokyo, Japan
      Namba, Kenryo, Tokyo, Japan
                                                                 <-- .
ΡĮ
      US 5958706
                             19990928
                                                               · <--
      WO 9735964 19971002
    ANSWER 24 OF 224 USPATFULL on STN
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DNA encoding a 2-acyltransferases

TI

Slabas, Antoni Ryszard, High Shincliffe, United Kingdom IN Brown, Adrian Paul, Shadforth, United Kingdom US 5945323 19990831 PIANSWER 25 OF 224 USPATFULL on STN L30 Microlocal calibration of digital printers ΤI Rao, Ravishankar, White Plains, NY, United States IN Thompson, Gerhard Robert, Wappingers Falls, NY, United States Tresser, Charles P., Mamaroneck, NY, United States Wu, Chai Wah, Ossining, NY, United States US 5943477 19990824 PΙ <--ANSWER 26 OF 224 USPATFULL on STN L30 ΤI Mutated penicillin G acylase genes Van Der Laan, Jan M., Breda, Netherlands IN Riemens, Adriana M., Delft, Netherlands Quax, Wilhelmus J., Voorschoten, Netherlands PΙ US 5891703 19990406 <--WO 9605318 19960222 L30ANSWER 27 OF 224 USPATFULL on STN Fusion protein-bound magnetic particles for recombinant production and TI magnetic separation of polypeptides of interest Matsunaga, Tadashi, Fuchu, Japan IN US 5861285 PΤ 19990119 < - -ANSWER 28 OF 224 CAPLUS COPYRIGHT 2005 ACS on STN T.30 Comparative genetics of capsular polysaccharide biosynthesis in ΤI Streptococcus pneumoniae types belonging to serogroup 19 SO Journal of Bacteriology (1999), 181(17), 5355-5364 CODEN: JOBAAY; ISSN: 0021-9193 L30 ANSWER 29 OF 224 CAPLUS COPYRIGHT 2005 ACS on STN The Caenorhabditis elegans unc-49 locus encodes multiple subunits of a TT heteromultimeric GABA receptor Journal of Neuroscience (1999), 19(13), 5348-5359 SO CODEN: JNRSDS; ISSN: 0270-6474 L30 ANSWER 30 OF 224 CAPLUS COPYRIGHT 2005 ACS on STN Analysis of the 5' portion of the type 19A capsule locus identifies two TI classes of cpsC, cpsD, and cpsE genes in Streptococcus pneumoniae Journal of Bacteriology (1999), 181(11), 3599-3605 so CODEN: JOBAAY; ISSN: 0021-9193 => d 130 31-224 Ti in so pn ANSWER 31 OF 224 CAPLUS COPYRIGHT 2005 ACS on STN Genome sequence of the radioresistant bacterium Deinococcus radiodurans R1 Science (Washington, D. C.) (1999), 286(5444), 1571-1577 SO CODEN: SCIEAS; ISSN: 0036-8075 ANSWER 32 OF 224 CAPLUS COPYRIGHT 2005 ACS on STN L30 Sequence and analysis of chromosome 4 of the plant Arabidopsis thaliana SO Nature (London) (1999), 402(6763), 769-777 CODEN: NATUAS; ISSN: 0028-0836 ANSWER 33 OF 224 CAPLUS COPYRIGHT 2005 ACS on STN L30 Sequence and analysis of chromosome 2 of the plant Arabidopsis thaliana Nature (London) (1999), 402 (6763), 760-768 SO CODEN: NATUAS; ISSN: 0028-0836

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- jelly: a polycystic kidney disease-like protein SO Human Molecular Genetics (1999), 8(3), 543-549 CODEN: HMGEE5; ISSN: 0964-6906
- L30 ANSWER 35 OF 224 CAPLUS COPYRIGHT 2005 ACS on STN
- TI The Genomic Organization and Polymorphism Analysis of the Human Niemann-Pick C1 Gene
- SO Biochemical and Biophysical Research Communications (1999), 261(2), 493-498
 CODEN: BBRCA9; ISSN: 0006-291X
- L30 ANSWER 36 OF 224 CAPLUS COPYRIGHT 2005 ACS on STN
- TI The functional role of alternative splicing of Ca2+-activated K+ channels in auditory hair cells
- SO Annals of the New York Academy of Sciences (1999), 868 (Molecular and Functional Diversity of Ion Channels and Receptors), 379-385 CODEN: ANYAA9; ISSN: 0077-8923
- L30 ANSWER 37 OF 224 CAPLUS COPYRIGHT 2005 ACS on STN
- TI Evidence for lateral gene transfer between Archaea and bacteria from genome sequence of Thermotoga maritima
- SO Nature (London) (1999), 399(6734), 323-329 CODEN: NATUAS; ISSN: 0028-0836
- L30 ANSWER 38 OF 224 CAPLUS COPYRIGHT 2005 ACS on STN
- TI Cloning, expression and electrophysiological characterization of glycine receptor alpha subunit from zebrafish
- SO Neuroscience (Oxford) (1999), 90(1), 303-317 CODEN: NRSCDN; ISSN: 0306-4522
- L30 ANSWER 39 OF 224 CAPLUS COPYRIGHT 2005 ACS on STN
- TI Identification of three novel members of the calcium-dependent chloride channel (CaCC) family predominantly expressed in the digestive tract and trachea
- SO FEBS Letters (1999), 455(3), 295-301 CODEN: FEBLAL; ISSN: 0014-5793
- L30 ANSWER 40 OF 224 CAPLUS COPYRIGHT 2005 ACS on STN
- TI The frost Gene of Neurospora crassa Is a Homolog of Yeast cdc1 and Affects Hyphal Branching via Manganese Homeostasis
- SO Fungal Genetics and Biology (1999), 28(3), 227-237 CODEN: FGBIFV; ISSN: 1087-1845
- L30 ANSWER 41 OF 224 CAPLUS COPYRIGHT 2005 ACS on STN
- TI Prediction of the coding sequences of unidentified human genes. XIV. The complete sequences of 100 new cDNA clones from brain which code for large proteins in vitro
- SO DNA Research (1999), 6(3), 197-205 CODEN: DARSE8; ISSN: 1340-2838
- L30 ANSWER 42 OF 224 CAPLUS COPYRIGHT 2005 ACS on STN
- TI The complete nucleotide sequence of RNA2 of blackcurrant reversion nepovirus
- SO Virus Research (1999), 65(1), 87-92 CODEN: VIREDF; ISSN: 0168-1702
- L30 ANSWER 43 OF 224 CAPLUS COPYRIGHT 2005 ACS on STN
- TI Identification of an inhibitory Zn2+ binding site on the human glycine receptor $\alpha 1$ subunit
- SO Journal of Physiology (Cambridge, United Kingdom) (1999), 520(1), 53-64
 CODEN: JPHYA7; ISSN: 0022-3751
- L30 ANSWER 44 OF 224 USPATFULL on STN

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Travis, Robert D., Tucson, AZ, United States
IN
              US 5810530
PΙ
                                                            19980922
                                                                                                                                       <--
         ANSWER 45 OF 224 CAPLUS COPYRIGHT 2005 ACS on STN DUPLICATE 3
L30
TI
          Animals cells expressing an insect GABA receptor subunit genes and
          screening for ligands of the receptor
IN
          Tomalski, Michael D.; Gant, Daniel B.
SO
          U.S., 28 pp.
          CODEN: USXXAM
          PATENT NO.
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PΙ
          US 5854002
                                                 Α
                                                               19981229
          ANSWER 46 OF 224 CAPLUS COPYRIGHT 2005 ACS on STN
L30
          Cloning, cDNA sequences, and antifungal and oxidase activities of plant
          antifungal proteins
IN
          Stuiver, Maarten Hendrik; Custers, Jerome Hubertus Henricus Victor;
          Sela-Buurlage, Marianne Beatrix; Melchers, Leo Sjoerd; Van
          Deventer-Troost, Johanna Pieternella Els; Lageweg, Wessel; Ponstein, Anne
          PCT Int. Appl., 139 pp.
SO
          CODEN: PIXXD2
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          AU 718274
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B1 20041117
A 19991027
A 20000327
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          NZ 334517
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A
          JP 2001502525
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          BR 9711291
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                                                 A1 20021114
          US 2002168735
L30 ANSWER 47 OF 224 CAPLUS COPYRIGHT 2005 ACS on STN
ΤI
          Mammalian cytokine, related reagents
          Knappe, Andrea; Fickenscher, Helmut; Fleckenstein, Bernard
IN
SO
          PCT Int. Appl., 63 pp.
          CODEN: PIXXD2
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L30 ANSWER 48 OF 224 CAPLUS COPYRIGHT 2005 ACS on STN
          Complete genome sequence of the methanogenic archaeon, Methanococcus
TI
          jannaschii
          Bult, Carol J.; White, Owen R.; Smith, Hamilton O.; Woese, Carl R.;
IN
        Venter, J. Craig
                                             the state of the s
          PCT Int. Appl., 615 pp.
SO
          CODEN: PIXXD2
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         US 6797466
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L30 ANSWER 49 OF 224 USPATFULL on STN
ΤI
             DNA encoding 2-acyltransferases
             Slabas, Antoni Ryszard, High Shincliffe, United Kingdom
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TТ

IN

Interference blind type bolt

Brown, Adrian Paul, Shadforth, United Kingdom

US 5843739 WO 9413814 19940623

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- The human glycine receptor subunit α3. GLRA3 gene structure, ΤI chromosomal localization, and functional characterization of alternative transcripts
- Journal of Biological Chemistry (1998), 273(31), 19708-19714 SO CODEN: JBCHA3; ISSN: 0021-9258
- ANSWER 51 OF 224 CAPLUS COPYRIGHT 2005 ACS on STN L30
- ΤI A transposition-induced mutant of Nostoc ellipsosporum implicates an arginine-biosynthetic gene in the formation of cyanophycin granules and of functional heterocysts and akinetes
- SO Microbiology (Reading, United Kingdom) (1998), 144(7), 1799-1805 CODEN: MROBEO; ISSN: 1350-0872
- ANSWER 52 OF 224 CAPLUS COPYRIGHT 2005 ACS on STN
- Rapid communication: nucleotide sequence of the coding region for the porcine β1-adrenergic receptor gene
- SO Journal of Animal Science (1998), 76(6), 1720-1721 CODEN: JANSAG; ISSN: 0021-8812
- L30 ANSWER 53 OF 224 CAPLUS COPYRIGHT 2005 ACS on STN
- P-type ATPases mediate sodium and potassium effluxes in Schwanniomyces occidentalis
- SO Journal of Biological Chemistry (1998), 273(3), 1640-1646 CODEN: JBCHA3; ISSN: 0021-9258
- ANSWER 54 OF 224 CAPLUS COPYRIGHT 2005 ACS on STN L30
- Molecular timing of primate divergences as estimated by two nonprimate calibration points
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- SO Journal of Molecular Evolution (1998), 46(4), 382-388 CODEN: JMEVAU; ISSN: 0022-2844
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- SO Journal of Neurobiology (1998), 37(2), 305-320 CODEN: JNEUBZ; ISSN: 0022-3034
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     Li, Yi; Kirkness, Ewen F.
SO
     U.S., 29 pp.
     CODEN: USXXAM
     PATENT NO.
                        KIND
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     CA 2212225
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                        A1
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                        A1
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L30 ANSWER 72 OF 224 CAPLUS COPYRIGHT 2005 ACS on STN
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ΤI
     transgenic plants with altered C14 sterol reductase levels
     Jang, Jyan-Chyun; Sheen, Jen
TN
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     PCT Int. Appl., 71 pp.
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    Proteins involved in the synthesis and assembly of the O-antigen of
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IN
    Lam, Joseph S.; Burrows, Lori; Charter, Deborah; De Kievit, Teresa
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     PCT Int. Appl., 194 pp.
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    PCT Int. Appl., 70 pp.
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- IN Cascieri, Margaret A.; Linemeyer, David L.; Macneil, Douglas J.; Shiao, Lin-Lin; Strader, Catherine; Weinberg, David H.; Tan, Carina P.
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- L30 ANSWER 223 OF 224 CAPLUS COPYRIGHT 2005 ACS on STN
- TI Complete nucleotide sequences of cloned copies of the RNA genes coding for the hemagglutinin and matrix proteins of a human influenza virus
- SO Developments in Cell Biology (Amsterdam) (1981), 7 (Replication Negat. Strand Viruses), 241-9
 CODEN: DCBIDD; ISSN: 0165-2265
- L30 ANSWER 224 OF 224 CAPLUS COPYRIGHT 2005 ACS on STN
- TI Cloning of influenza cDNA into M13: the sequence of the RNA segment

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encoding the A/PR/8/34 matrix protein
SO
     Nucleic Acids Research (1980), 8(9), 1965-74
     CODEN: NARHAD; ISSN: 0305-1048
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L21
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=> FIL REGISTRY
COST IN U.S. DOLLARS

L29

L30 L31

L32

0 S L28 AND DUP REM

1 S L30 AND (VII)

224 DUP REM L28 (7 DUPLICATES REMOVED)

0 S L30 AND (FACTOR (W) VII)

SINCE FILE TOTAL ENTRY SESSION

FULL ESTIMATED COST 470.73 932.54

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS) SINCE FILE TOTAL

ENTRY

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Property values tagged with IC are from the ZIC/VINITI data file provided by InfoChem.

STRUCTURE FILE UPDATES: 3 JAN 2005 HIGHEST RN 807382-78-1 DICTIONARY FILE UPDATES: 3 JAN 2005 HIGHEST RN 807382-78-1

TSCA INFORMATION NOW CURRENT THROUGH MAY 21, 2004

Please note that search-term pricing does apply when conducting SmartSELECT searches.

Crossover limits have been increased. See HELP CROSSOVER for details.

Experimental and calculated property data are now available. For more information enter HELP PROP at an arrow prompt in the file or refer to the file summary sheet on the web at: http://www.cas.org/ONLINE/DBSS/registryss.html

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=> FIL CAPLUS BIOSIS MEDLINE PCTFULL USPATFULL EUROPATFULL JAPIO SCISEARCH EMBASE COST IN U.S. DOLLARS SINCE FILE TOTAL

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FILE 'PCTFULL' ENTERED AT 15:46:57 ON 04 JAN 2005 COPYRIGHT (C) 2005 Univentio

FILE 'USPATFULL' ENTERED AT 15:46:57 ON 04 JAN 2005
CA INDEXING COPYRIGHT (C) 2005 AMERICAN CHEMICAL SOCIETY (ACS)

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'EUROPATFULL' IS NOT A VALID FILE NAME
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L25

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L5
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     FILE 'REGISTRY' ENTERED AT 15:17:27 ON 04 JAN 2005
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L17
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L18
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L19
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L21
L22
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FILE 'CAPLUS, BIOSIS, MEDLINE, PCTFULL, USPATFULL, JAPIO, SCISEARCH,

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	EMBASE' ENTERED AT 15:31:55 ON 04 JAN 2005					
L28	231 S L26 AND PY<=1999					
L29	0 S L28 AND DUP REM					
L30	224 DUP REM L28 (7 DUPLICATES REMOVED)					
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L33 L34	FILE 'REGISTRY' ENTERED AT 15:44:46 ON 04 JAN 2005 0 S [WFL] . [VI] [LIMV] [WFLM] . [W] [-P] [RKLW]/SQSP AND SQL=18 4 S [WFL] . [VI] [LIMV] [WFLM] . [W] [-P] [RKLW]/SQSP AND SQL<30					
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